

Nobivac DHPPi Vet. lyophilisate and solvent for suspension for injection for dogs

Authorised

- Canine parainfluenza virus, strain Cornell, Live
- Canine adenovirus 2, strain Manhattan LPV3, Live
- Canine distemper virus, strain Onderstepoort, Live
- Canine parvovirus, strain 154, Live

Product identification

Medicine name:

Nobivac DHPPi Vet. lyophilisate and solvent for suspension for injection for dogs

Active substance:

Canine parainfluenza virus, strain Cornell, Live
Canine adenovirus 2, strain Manhattan LPV3, Live
Canine distemper virus, strain Onderstepoort, Live
Canine parvovirus, strain 154, Live

Target species:

Dog

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Canine parainfluenza virus, strain Cornell, Live

316228.00 50% tissue culture infectious dose / 1.00 millilitre(s)

Canine adenovirus 2, strain Manhattan LPV3, Live

10000.00 50% tissue culture infectious dose / 1.00 millilitre(s)

Canine distemper virus, strain Onderstepoort, Live

10000.00 50% tissue culture infectious dose / 1.00 millilitre(s)

Canine parvovirus, strain 154, Live

10000000.00 50% tissue culture infectious dose / 1.00 millilitre(s)

Pharmaceutical form:

Lyophilisate and solvent for suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07AD04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Denmark

Available in:

Denmark

Package description:

10 x 1 dose vials in a carton or plastic box

25 x 1 dose vials in a carton or plastic box

50 x 1 dose vials in carton or plastic box

5 x 1 dose vials in a carton or plastic box

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Intervet International B.V.

Marketing authorisation date:

6/12/2000

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

Danish Medicines Agency

Authorisation number:

19406

Date of authorisation status change:

6/12/2000

Reference member state:

Denmark

Procedure number:

DK/V/0103/001

Concerned member states:

Norway Slovenia

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

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PI.pdf